

Amendment to the Claims

This listing of claims will replace all prior versions and listings of claims in the above-referenced application.

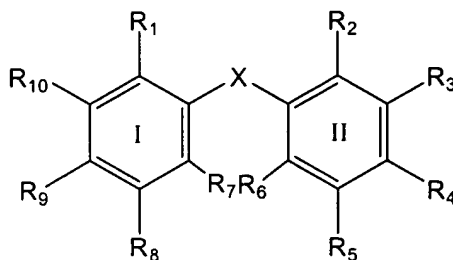
1-50. (canceled)

51. (new) A method of treating a subject in need of treatment for malaria, wherein the subject is infected with a malaria parasite, the method comprising the step of:

administering an antimalarial composition comprising a compound that is an inhibitor of fatty acid synthesis in the malaria parasite to the subject.

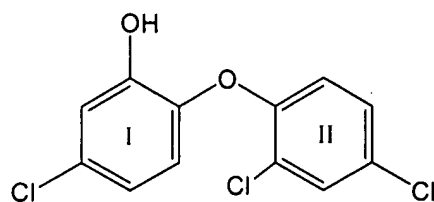
52. (new) The method of claim 51, wherein the inhibitor of fatty acid synthesis is a hydroxydiphenyl ether.

53. (new) The method of claim 52, wherein the hydroxydiphenyl ether has general formula 2 given below wherein the two phenyl rings (I & II) are joined by an oxygen atom ($X=O$) and either R_1 or R_2 represents a hydroxy (OH) group with the other being a hydrogen atom, respectively, or both being hydroxy groups, and wherein R_3 to R_{10} of the phenyl rings I and II are selected from the group consisting of: chlorine, bromine, iodine, hydrogen, hydroxyl groups, aldehyde groups, keto groups, and ester groups.



Formula 2

54. (new) The method of claim 52, wherein the inhibitor of fatty acid synthesis is triclosan having formula 1 given below:



Formula 1

55. (new) The method of claim 52, wherein the composition further comprises one or more known antimalarial agents and a pharmaceutically acceptable adjuvant, diluent, or carrier.
56. (new) The method of claim 55, wherein the known antimalarial agent is selected from the group consisting of: quinine, atabrine, chloroquine, mefloquine, primaquine, anti-folates, artemisinin, artemether, and artesunate.
57. (new) The method of claim 52, wherein the composition is administered by injection.
58. (new) The method of claim 52, wherein the amount of the inhibitor of fatty acid synthesis administered is in the dosage range of 0.03 mg/kg to 100 mg/kg.
59. (new) The method of claim 52, wherein the compound inhibits FabI (enoyl ACP reductase) in the malaria parasite.
60. (new) The method of claim 51, wherein the composition further comprises one or more known antimalarial agents and a pharmaceutically acceptable adjuvant, diluent, or carrier.
61. (new) The method of claim 60, wherein the known antimalarial agent is selected from the group consisting of: quinine, atabrine, chloroquine, mefloquine, primaquine, anti-folates, artemisinin, artemether, and artesunate.
62. (new) The method of claim 51, wherein the inhibitor of fatty acid synthesis in the malaria parasite is an inhibitor of FabI (enoyl-ACP reductase).
63. (new) The method of claim 51, wherein the malaria parasite is *P. falciparum*.